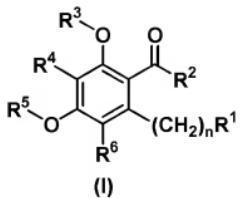


a.) Amendment to the Claims

1. (Currently Amended) A Hsp90 family protein inhibitor comprising, as an active ingredient, A method of inhibiting a Hsp90 family protein, comprising administering to a patient in need thereof, an effective amount of a benzoyl compound represented by general formula (I):



[wherein

n represents an integer of 0 to 10;

R¹ represents a hydrogen atom, hydroxy, cyano, carboxy, nitro, halogen, substituted or unsubstituted lower alkyl, substituted or unsubstituted lower alkenyl, substituted or unsubstituted lower alkynyl, substituted or unsubstituted lower alkoxy, substituted or unsubstituted cycloalkyl, substituted or unsubstituted lower alkoxy carbonyl, substituted or unsubstituted lower alkanoyloxy, substituted or unsubstituted heterocyclic alkyl, substituted or unsubstituted aryl, substituted or unsubstituted arylsulfonyl, a substituted or unsubstituted heterocyclic group, CONR⁷R⁸ (wherein R⁷ and R⁸ independently represent a hydrogen atom, substituted or unsubstituted lower alkyl, substituted or unsubstituted cycloalkyl, substituted or unsubstituted lower alkanoyl,

substituted or unsubstituted aryl, a substituted or unsubstituted heterocyclic group, substituted or unsubstituted aralkyl, substituted or unsubstituted heterocyclic alkyl, or substituted or unsubstituted aroyl, or R⁷ and R⁸ form a substituted or unsubstituted heterocyclic group together with the adjacent nitrogen atom) or NR⁹R¹⁰ (wherein R⁹ and R¹⁰ have the same meanings as the above R⁷ and R⁸, respectively);

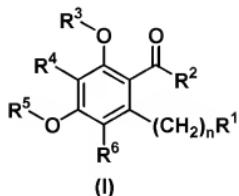
R² represents substituted or unsubstituted lower alkyl, substituted or unsubstituted lower alkenyl, substituted or unsubstituted lower alkynyl, substituted or unsubstituted cycloalkyl, substituted or unsubstituted aryl, or a substituted or unsubstituted heterocyclic group;

R³ and R⁵ independently represent a hydrogen atom, substituted or unsubstituted lower alkyl, substituted or unsubstituted lower alkenyl, substituted or unsubstituted lower alkanoyl, substituted or unsubstituted cycloalkyl, substituted or unsubstituted aralkyl, or substituted or unsubstituted aroyl; and

R⁴ and R⁶ independently represent a hydrogen atom, hydroxy, halogen, cyano, nitro, substituted or unsubstituted lower alkyl, substituted or unsubstituted lower alkenyl, substituted or unsubstituted lower alkynyl, substituted or unsubstituted lower alkoxy, substituted or unsubstituted cycloalkyl, amino, lower alkylamino, di-lower alkylamino, carboxy, substituted or unsubstituted lower alkoxy carbonyl, substituted or unsubstituted lower alkanoyl, substituted or unsubstituted aryloxy, substituted or unsubstituted aryl, a substituted or unsubstituted heterocyclic group, substituted or unsubstituted aralkyl, or substituted or unsubstituted heterocyclic alkyl], or

a prodrug thereof, or a pharmaceutically acceptable salt of said benzoyl compound or said prodrug.

2. (Currently Amended) An Hsp90 family protein inhibitor comprising, as an active ingredient, A method of inhibiting a Hsp90 family protein, comprising administering to a patient in need thereof, an effective amount of a benzoyl compound represented by general formula (I):



[wherein

n represents an integer of 0 to 10;

R¹ represents a hydrogen atom, hydroxy, cyano, carboxy, nitro, halogen, substituted or unsubstituted lower alkyl, substituted or unsubstituted lower alkenyl, substituted or unsubstituted lower alkynyl, substituted or unsubstituted lower alkoxy, substituted or unsubstituted cycloalkyl, substituted or unsubstituted lower alkoxy carbonyl, substituted or unsubstituted lower alkanoyloxy, substituted or unsubstituted heterocyclic alkyl, substituted or unsubstituted aryl, substituted or unsubstituted arylsulfonyl, a substituted or unsubstituted heterocyclic group, CONR⁷R⁸ (wherein R⁷ and R⁸

independently represent a hydrogen atom, substituted or unsubstituted lower alkyl, substituted or unsubstituted cycloalkyl, substituted or unsubstituted lower alkanoyl, substituted or unsubstituted aryl, a substituted or unsubstituted heterocyclic group, substituted or unsubstituted aralkyl, substituted or unsubstituted heterocyclic alkyl, or substituted or unsubstituted aroyl, or R⁷ and R⁸ form a substituted or unsubstituted heterocyclic group together with the adjacent nitrogen atom) or NR⁹R¹⁰ (wherein R⁹ and R¹⁰ have the same meanings as the above R⁷ and R⁸, respectively);

R² represents substituted or unsubstituted lower alkyl, substituted or unsubstituted lower alkenyl, substituted or unsubstituted lower alkynyl, substituted or unsubstituted cycloalkyl, substituted or unsubstituted aryl, or a substituted or unsubstituted heterocyclic group;

R³ and R⁵ independently represent a hydrogen atom, substituted or unsubstituted lower alkyl, substituted or unsubstituted lower alkenyl, substituted or unsubstituted lower alkanoyl, substituted or unsubstituted cycloalkyl, substituted or unsubstituted aralkyl, or substituted or unsubstituted aroyl; and

R⁴ and R⁶ independently represent a hydrogen atom, hydroxy, halogen, cyano, nitro, substituted or unsubstituted lower alkyl, substituted or unsubstituted lower alkenyl, substituted or unsubstituted lower alkynyl, substituted or unsubstituted lower alkoxy, substituted or unsubstituted cycloalkyl, amino, lower alkylamino, di-lower alkylamino, carboxy, substituted or unsubstituted lower alkoxy carbonyl, substituted or unsubstituted lower alkanoyl, substituted or unsubstituted aryloxy, substituted or

unsubstituted aryl, a substituted or unsubstituted heterocyclic group, substituted or unsubstituted aralkyl, or substituted or unsubstituted heterocyclic alkyl],

or a pharmaceutically acceptable salt thereof.

3. (Currently Amended) The Hsp90 family protein inhibitor The method according to claim 2, wherein R¹ is a hydrogen atom, hydroxy, cyano, carboxy, nitro, halogen, substituted or unsubstituted lower alkyl, substituted or unsubstituted lower alkynyl, substituted or unsubstituted lower alkoxy, substituted or unsubstituted cycloalkyl, substituted or unsubstituted lower alkoxy carbonyl, substituted or unsubstituted lower alkanoyloxy, substituted or unsubstituted heterocyclic alkyl, substituted or unsubstituted aryl, substituted or unsubstituted arylsulfonyl, CONR⁷R⁸ or NR⁹R¹⁰.

4. (Currently Amended) The Hsp90 family protein inhibitor The method according to claim 2, wherein R¹ is substituted or unsubstituted lower alkyl, substituted or unsubstituted lower alkynyl, substituted or unsubstituted lower alkoxy, substituted or unsubstituted cycloalkyl, substituted or unsubstituted lower alkoxy carbonyl, substituted or unsubstituted heterocyclic alkyl, substituted or unsubstituted aryl, CONR⁷R⁸ or NR⁹R¹⁰.

5. (Currently Amended) The Hsp90 family protein inhibitor The method according to claim 2, wherein R¹ is CONR⁷R⁸.

6. (Currently Amended) The Hsp90 family protein inhibitor The method according to any one of claims 2 to 5, wherein R² is substituted or unsubstituted aryl or a substituted or unsubstituted aromatic heterocyclic group.

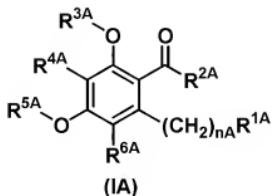
7. (Currently Amended) The Hsp90 family protein inhibitor The method according to claim 6, wherein R⁴ is a hydrogen atom, hydroxy or halogen.

8. (Currently Amended) The Hsp90 family protein inhibitor The method according to claim 7, wherein R³ and R⁵, which may be the same or different, each are a hydrogen atom, substituted or unsubstituted lower alkyl, substituted or unsubstituted lower alkenyl, substituted or unsubstituted lower alkanoyl, or substituted or unsubstituted aroyl.

9. (Currently Amended) The Hsp90 family protein inhibitor The method according to claim 6, wherein R³, R⁴ and R⁵ each are a hydrogen atom.

10. (Currently Amended) The Hsp90 family protein inhibitor The
method according to claim 9, wherein R⁶ is a hydrogen atom, lower alkyl, halogen or aryl.

11. (Previously Presented) A benzoyl compound represented by general formula (IA):



[wherein

nA represents an integer of 1 to 5;

R^{1A} represents substituted or unsubstituted lower alkyl, substituted or unsubstituted lower alkoxy, substituted or unsubstituted cycloalkyl, substituted or unsubstituted lower alkoxy carbonyl, substituted or unsubstituted heterocyclic alkyl, substituted or unsubstituted aryl, CONR⁷R⁸ (wherein R⁷ and R⁸ independently represent a hydrogen atom, substituted or unsubstituted lower alkyl, substituted or unsubstituted cycloalkyl, substituted or unsubstituted lower alkanoyl, substituted or unsubstituted aryl, a substituted or unsubstituted heterocyclic group, substituted or unsubstituted aralkyl, substituted or unsubstituted heterocyclic alkyl, or substituted or unsubstituted aroyl, or R⁷ and R⁸ form a substituted or unsubstituted heterocyclic group together with the adjacent

nitrogen atom) or NR⁹R¹⁰ (wherein R⁹ and R¹⁰ have the same meanings as the above R⁷ and R⁸, respectively);

R^{2A} represents substituted or unsubstituted aryl or a substituted or unsubstituted aromatic heterocyclic group;

R^{3A} and R^{5A} independently represent a hydrogen atom, substituted or unsubstituted lower alkyl, substituted or unsubstituted lower alkenyl, substituted or unsubstituted lower alkanoyl, substituted or unsubstituted cycloalkyl, substituted or unsubstituted aralkyl, or substituted or unsubstituted aroyl;

R^{4A} represents a hydrogen atom, hydroxy or halogen; and

R^{6A} represents a hydrogen atom, halogen, cyano, nitro, substituted or unsubstituted lower alkyl, substituted or unsubstituted lower alkenyl, substituted or unsubstituted lower alkynyl, substituted or unsubstituted lower alkoxy, substituted or unsubstituted cycloalkyl, amino, lower alkylamino, di-lower alkylamino, carboxy, substituted or unsubstituted lower alkoxy carbonyl, substituted or unsubstituted lower alkanoyl, substituted or unsubstituted aryloxy, substituted or unsubstituted aryl, a substituted or unsubstituted heterocyclic group, substituted or unsubstituted aralkyl, or substituted or unsubstituted heterocyclic alkyl; provided that:

(i) when R^{3A} and R^{5A} each are methyl, R^{4A} and R^{6A} each are a hydrogen atom, and

-(CH₂)_{nA}R^{1A} is

(a) methoxycarbonylmethyl,

R^2A is not a group selected from the group consisting of 2,4,6-trimethoxy-5-methoxycarbonyl-3-nitrophenyl, 3-cyano-2,4,6-trimethoxyphenyl, 5-cyano-2-ethoxy-4,6-dimethoxy-3-nitrophenyl, 2,6-dimethoxyphenyl, 2-chloro-6-methoxyphenyl and 2-chloro-4,6-dimethoxy-5-methoxycarbonyl-3-nitrophenyl,

(b) ethoxycarbonylmethyl;

R^2A is not 2,4,6-trimethoxy-3-methoxycarbonyl-phenyl, and

(c) N,N-dimethylaminomethyl,

R^2A is not phenyl;

(ii) when R^3A , R^4A , R^5A and R^6A each are a hydrogen atom, and -

$(CH_2)_nAR^1A$ is

(a) 2-(acetoxymethyl)heptyl, 3-oxopentyl or pentyl,

R^2A is not 6-hydroxy-4-methoxy-3-methoxycarbonyl-2-pentylphenyl,

(b) 3-oxopentyl,

R^2A is not a group selected from the group consisting of 3-benzyloxycarbonyl-6-hydroxy-4-methoxy-2-pentylphenyl and 3-carboxy-6-hydroxy-4-methoxy-2-pentylphenyl, and

(c) n-propyl,

R^2A is not 2,4-dihydroxy-6-[(4-hydroxy-2-oxopyran-6-yl)methyl]phenyl;

(iii) when R^3A and R^4A each are a hydrogen atom, R^5A is methyl, R^6A is methoxycarbonyl, and $-(CH_2)_nAR^1A$ is pentyl;

R^2A is not a group selected from the group consisting of 6-[2-(acetoxymethyl)heptyl]-2,4-dihydroxyphenyl, 2,4-dihydroxy-6-pentylphenyl and 2,4-dihydroxy-6-(3-oxopentyl)phenyl;

(iv) when R^3A and R^5A each are benzyl, R^4A and R^6A each are a hydrogen atom, and $-(CH_2)_nAR^1A$ is 3-oxopentyl,

R^2A is not a group selected from the group consisting of 6-benzyloxy-4-methoxy-3-methoxycarbonyl-2-pentylphenyl and 6-benzyloxy-3-benzyloxycarbonyl-4-methoxy-2-pentylphenyl;

(v) when R^3A is benzyl, R^4A is a hydrogen atom, R^5A is methyl, $-(CH_2)_nAR^1A$ is pentyl, and R^6A is methoxycarbonyl or benzyloxycarbonyl,

R^2A is not 2,4-bis(benzyloxy)-6-(3-oxopentyl)-phenyl;

(vi) when R^3A and R^4A each are a hydrogen atom, R^5A is methyl, $-(CH_2)_nAR^1A$ is pentyl, and R^6A is carboxy or benzyloxycarbonyl,

R^2A is not 2,4-dihydroxy-6-(3-oxopentyl)phenyl; and

(vii) when R³A, R⁴A and R⁶A each are a hydrogen atom, R⁵A is n-propyl, and -(CH₂)_nAR¹A is 5-(1,1-dimethylpropyl)-4-(2-hydrobenzotriazol-2-yl)-2-hydroxyphenylmethyl,

R²A is not phenyl],

or a pharmaceutically acceptable salt thereof.

12. (Original) The benzoyl compound according to claim 11, wherein R²A is a substituted or unsubstituted aromatic heterocyclic group, substituted aryl having 1 to 3 substituents, or aryl, or a pharmaceutically acceptable salt thereof.

13. (Previously Presented) The benzoyl compound according to claim 12, wherein R³A and R⁵A are independently a hydrogen atom, substituted or unsubstituted lower alkyl, substituted or unsubstituted lower alkanoyl, substituted or unsubstituted aroyl, or substituted or unsubstituted lower alkenyl, or a pharmaceutically acceptable salt thereof.

14. (Previously Presented) The benzoyl compound according to claim 12, wherein R³A, R⁴A and R⁵A each are a hydrogen atom, or a pharmaceutically acceptable salt thereof.

15. (Previously Presented) The benzoyl compound according to any one of claims 11 to 14, wherein R^{1A} is CONR⁷R⁸, or a pharmaceutically acceptable salt thereof.

16. (Previously Presented) The benzoyl compound according to any one of claims 11 to 14, wherein R^{6A} is a hydrogen atom, lower alkyl, halogen or aryl, or a pharmaceutically acceptable salt thereof.

17. (Previously Presented) A pharmaceutical composition comprising, as an active ingredient, the benzoyl compound according to claim 16 or a pharmaceutically acceptable salt of said benzoyl compound, together with a pharmaceutically acceptable carrier.

18. (Previously Presented) A pharmaceutical composition comprising, as an active ingredient, a prodrug of the benzoyl compound according to claim 16 or a pharmaceutically acceptable salt thereof, together with a pharmaceutically acceptable carrier.

19. (Previously Presented) A pharmaceutical composition comprising, as an active ingredient, the benzoyl compound according to any one of claims 11 to 14, or

a pharmaceutically acceptable salt of said benzoyl compound, together with a pharmaceutically acceptable carrier.

20. (Previously Presented) A pharmaceutical composition comprising, as an active ingredient, a prodrug of the benzoyl compound according to any one of claims 11 to 14 or a pharmaceutically acceptable salt thereof, together with a pharmaceutically acceptable carrier.

Claims 21-25 (Cancelled).

26. (Previously Presented) A method of treating a disease associated with an Hsp90 family protein or a Hsp90 client protein, which comprises administering an effective amount of the benzoyl compound according to any one of claims 11 to 14 or a prodrug thereof, or a pharmaceutically acceptable salt of said benzoyl compound or said prodrug.

27. (Previously Presented) A method of treating malignant tumors, which comprises administering an effective amount of the benzoyl compound according to any one of claims 11 to 14 or a prodrug thereof, or a pharmaceutically acceptable salt of said benzoyl compound or said prodrug.

Claims 28-31 (Cancelled).